GREENEX HAND SANITIZER- ethanol liquid
Cleanslate Group LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Greenex Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on the skin after changing diapers, assisting ill persons, or before contact with a person under medical care or treatment.
- Recommended for repeated use.

Warnings

- For external use only.
- Keep out of eyes, ears, or mouth.
- Discontinue use if irritation occurs.
- Keep out of reach of children.

FLAMMABLE, KEEP AWAY FROM FIRE OR FLAME.

Directions

- If hands are visibly soiled, wash with soap and water and dry hands.
- Wet hands thoroughly with product, especially the area under the fingernails and allow to dry without rinsing.

Inactive Ingredients benzalkonium chloride, chlorhexidine gluconate, isopropyl alcohol, PEG 10 dimethicone, PEG-14M, phenoxyethanol, polyquaternium 10, water.

GREENEX

2 Bergen Turnpike

Ridgefield Park, NJ 07660

MADE IN THE USA

GREENEX

HAND SANITIZER

Kills more than 99.99% germs in 15 seconds

1000 ml

GREENEX HAND SANITIZER

ethanol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80586-202
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
POLYETHYLENE OXIDE 600000 (UNII: 2126FD486L)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYQUATERNIUM-10 (1000 MPA.S AT 2%) (UNII: GMR4PEN8PK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80586-202-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/13/2021	

Labeler - Cleanslate Group LLC (117657934)

Revised: 4/2021 Cleanslate Group LLC